

TRANSMITTAL LETTER TO THE UNITED STATES
DESIGNATED/ELECTED OFFICE (DO/EO/US)
CONCERNING A FILING UNDER 35 U.S.C. 371412 Rec'd PCT/PTO 13 MAR 2000
365-442P

U.S. APPLICATION NO. (If known, see 37 CFR 1.5)

09/508405

INTERNATIONAL APPLICATION NO.

PCT/FI98/00764

INTERNATIONAL FILING DATE

September 28, 1998

PRIORITY DATE CLAIMED

September 26, 1997

TITLE OF INVENTION

HIGH MELT STRENGTH POLYPROPYLENE

APPLICANT(S) FOR DO/EO/US

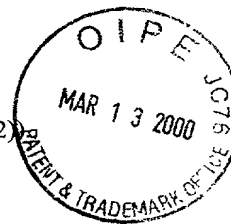
HUOVINEN, Paivi; HARLIN, Ali; JAASKELAINEN, Pirjo; KARBASI, Amir; MANNER, Nina

Applicant herewith submits to the United States Designated/Elected Office (DO/EO/US) the following items and other information:

1. ☒ This is a **FIRST** submission of items concerning a filing under 35 U.S.C. 371.
2. ☐ This is a **SECOND** or **SUBSEQUENT** submission of items concerning a filing under 35 U.S.C. 371.
3. ☒ This express request to begin national examination procedures (35 U.S.C. 371(f)) at any time rather than delay examination until the expiration of the applicable time limit set in 35 U.S.C. 371(b) and PCT Articles 22 and 39 (1).
4. ☒ A proper Demand for International Preliminary Examination was made by the 19th month from the earliest claimed priority date
5. ☒ A copy of the International Application as filed (35 U.S.C. 371(c)(2))
- a. ☒ is transmitted herewith (required only if not transmitted by the International Bureau). WO 99/16797
- b. ☒ has been transmitted by the International Bureau.
- c. ☐ is not required, as the application was filed in the United States Receiving Office (RO/US).
6. ☐ A translation of the International Application into English (35 U.S.C. 371(c)(3)).
7. ☒ Amendments to the claims of the International Application under PCT Article 19 (35 U.S.C. 371(c)(2))
- a. ☐ are transmitted herewith (required only if not transmitted by the International Bureau).
- b. ☐ have been transmitted by the International Bureau.
- c. ☐ have not been made; however, the time limit for making such amendments has NOT expired.
- d. ☒ have not been made and will not be made.
8. ☐ A translation of the amendments to the claims under PCT Article 19 (35 U.S.C. 371(c)(3)).
9. ☐ An oath or declaration of the inventor(s) (35 U.S.C. 371(c)(4)).
10. ☐ A translation of the annexes to the International Preliminary Examination Report under PCT Article 36 (35 U.S.C. 371(c)(5)).

Items 11. to 16. below concern document(s) or information included:

11. ☒ An Information Disclosure Statement under 37 CFR 1.97 and 1.98.-1449 and International Search Report (PCT/ISA/210)
12. ☐ An assignment document for recording. A separate cover sheet in compliance with 37 CFR 3.28 and 3.31 is included.
13. ☒ A **FIRST** preliminary amendment.
☐ A **SECOND** or **SUBSEQUENT** preliminary amendment.
14. ☐ A substitute specification.
15. ☐ A change of power of attorney and/or address letter.
16. ☒ Other items or information:
1.) PCT Request (PCT/RO/101)
2.) International Preliminary Examination Report (PCT/IPEA/409)
3.) Four (4) sheets of Formal Drawings



09/508405

PCT/FI98/00764

365-442P

17. ☒ The following fees are submitted:**BASIC NATIONAL FEE (37 CFR 1.492(a)(1)-(5):**

Neither international preliminary examination fee (37 CFR 1.482)
nor international search fee (37 CFR 1.445(a)(2)) paid to USPTO
and International Search Report not prepared by the EPO or JPO. \$970.00

International preliminary examination fee (37 CFR 1.482) not paid to
USPTO but International Search Report prepared by the EPO or JPO \$840.00

International preliminary examination fee (37 CFR 1.482) not paid to USPTO
but international search fee (37 CFR 1.445(a)(2)) paid to USPTO. \$690.00

International preliminary examination fee (37 CFR 1.482) paid to USPTO
but all claims did not satisfy provisions of PCT Article 33(1)-(4) \$670.00

International preliminary examination fee (37 CFR 1.482) paid to USPTO
and all claims satisfied provisions of PCT Article 33(1)-(4). \$96.00

ENTER APPROPRIATE BASIC FEE AMOUNT =

Surcharge of \$130.00 for furnishing the oath or declaration later than ☐ 20 ☒ 30
months from the earliest claimed priority date (37 CFR 1.492(e)).

CLAIMS	NUMBER FILED	NUMBER EXTRA	RATE
Total Claims	51 - 20 =	31	X \$18.00
Independent Claims	2 - 3 =	0	X \$78.00
MULTIPLE DEPENDENT CLAIM(S) (if applicable) Yes			+ \$260.00

TOTAL OF ABOVE CALCULATIONS =

Reduction of 1/2 for filing by small entity, if applicable. Verified Small Entity statement
must also be filed (Note 37 CFR 1.9, 1.27, 1.28).

SUBTOTAL =

Processing fee of \$130.00 for furnishing the English translation later than ☐ 20 ☐ 30
months from the earliest claimed priority date (37 CFR 1.492(f)).

TOTAL NATIONAL FEE =

Fee for recording the enclosed assignment (37 CFR 1.21(h)). The assignment must be
accompanied by an appropriate cover sheet (37 CFR 3.28, 3.31). \$40.00 per property +

TOTAL FEES ENCLOSED =**CALCULATIONS PTO USE ONLY**

514 Rec'd PCT/PTO 13 MAR 2000

\$ 970.00

\$ 130.00

\$ 558.00

\$ 0

\$ 260.00

\$ 1,918.00

\$ 0

\$ 1,918.00

\$ 0

\$ 1,918.00

\$ 0

\$ 1,918.00

Amount to be:
refunded \$

charged \$

a. ☒ A check in the amount of \$ 1,918.00 to cover the above fees is enclosed.

b. ☐ Please charge my Deposit Account. No. _____ in the amount of \$ _____ to cover the above fees.
A duplicate copy of this sheet is enclosed.

c. ☒ The Commissioner is hereby authorized to charge any additional fees which may be required, or credit any
overpayment to Deposit Account No. 02-2448.

NOTE: Where an appropriate time limit under 37 CFR 1.494 or 1.495 has not been met, a petition to revive (37 CFR 1.137(a) or (b)) must be filed and granted to restore the application to pending status.

Send all correspondence to:

Birch, Stewart, Kolasch & Birch, LLP or Customer No. 2292
P.O. Box 747
Falls Church, VA 22040-0747
(703)205-8000

SIGNATURE

SVENSSON, LEONARD R.
NAME

#30,330 (LRS)
REGISTRATION NUMBER

/cqc

09/508405
514 Rec'd PCT/PTO 13 MAR 2000

PATENT
365-442P

IN THE U.S. PATENT AND TRADEMARK OFFICE

Applicant: HUOVINEN, Paivi et al.
Int'l. Appl. No.: PCT/FI98/00764
Appl. No.: New Group:
Filed: March 13, 2000 Examiner:
For: HIGH MELT STRENGTH POLYPROPYLENE

PRELIMINARY AMENDMENT

BOX PATENT APPLICATION

Assistant Commissioner for Patents
Washington, DC 20231

March 13, 2000

Sir:

The following Preliminary Amendments and Remarks are respectfully submitted in connection with the above-identified application.

AMENDMENTS

IN THE SPECIFICATION:

Please amend the specification as follows:

Before line 1, insert --This application is the national phase under 35 U.S.C. § 371 of PCT International Application No. PCT/FI98/00764 which has an International filing date of September 28, 1998, which designated the United States of America.--

IN THE CLAIMS:

Please amend the claims as follows:

Claim 4: Line 1, change "any of the claims 1 to 3" to
--claim 1--

Claim 5: Lines 1 to 2, change "any of the claims 1 to 4"
to --claim 1--

Claim 11: Line 1, change "any one of the claims 8 to 10"
to --claim 8--

Claim 13: Line 1, change "any of the claims 9 to 10" to
--claim 9--

Claim 17: Line 1, change "any one of the preceding claims"
to --claim 1--

Claim 18: Line 1, change "any one of the preceding claims"
to --claim 1--

Claim 21: Line 1, change "any one of the preceding claims"
to --claim 1--

Claim 22: Line 1, change "any one of the preceding claims"
to --claim 1--

Claim 27: Line 1, delete "or 26"

Claim 28: Line 1, change "any one of the preceding claims" to -
-claim 1--

Claim 39: Line 1, change "any of claims 8 to 13" to --claim 8--

Claim 41: Line 1, change "any one of the preceding claims" to -
-claim 1--

Claim 42: Line 1, change "any of claims 1 to 4" to --claim 1--;
and

Line 2, change "any of claims 5 to 7" to --claim 5--

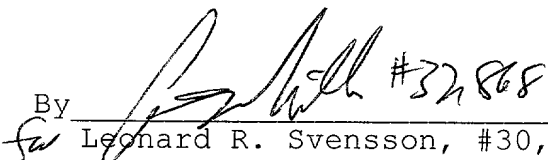
REMARKS

The specification has been amended to provide a cross-reference to the previously filed International Application. The claims have also been amended to delete the improper multiple dependencies. Favorable action on the above-identified application is respectfully requested.

If necessary, the Commissioner is hereby authorized in this, concurrent, and future replies, to charge payment or credit any overpayment to Deposit Account No. 02-2448 for any additional fees required under 37 C.F.R. § 1.16 or under 37 C.F.R. § 1.17; particularly, extension of time fees.

Respectfully submitted,

BIRCH, STEWART, KOLASCH & BIRCH, LLP

By  #32868
for Leonard R. Svensson, #30,330

LRS/cqc
365-442P

P.O. Box 747
Falls Church, VA 22040-0747
(703) 205-8000

(Rev. 01/08/2000)

4/PRTS

09/508405
514 Rec'd PCT/PTO 13 MAR 2000

1

HIGH MELT STRENGTH POLYPROPYLENE

Background of the Invention

5 Field of the Invention

10 The present invention relates to high melt strength polymers. In particular, the present invention concerns linear poly-propylene having high melt strength and comprising a high molecular weight portion and a low or medium molecular weight portion. The invention further concerns a process for preparing linear homopolymers and copolymers of propylene having high melt strength. Finally, the present invention relates to the use of the new polypropylene for the manufacture of products by, e.g., foaming, melt-stage thermo-forming, blow moulding, extrusion coating, fibre forming and injection moulding.

15 Description of Related Art

20 An important characteristic of polypropylene (PP) homopolymers is their stiffness, which is closely related to the crystallinity and the molecular structure of the polymer chain. Further, PP homopolymers have good resistance to deformation at elevated temperatures, high tensile strength, surface hardness, and good toughness at ambient temperatures. By copolymerizing propylene with other monomers it is possible further to improve the impact resistance.

25 The thermoformability of polypropylene in melt phase is, however, rather poor, which can be mentioned as a problem with polypropylene. This feature is primarily due to sheet sag during heating and forming processes. Generally, a material which is thermo-formable in the melt-phase should possess a viscous component to allow for flow when stressed and a sufficiently elastic component to resist flow when not stressed. Another important factor affecting thermoforming is the melt drawability of the polymer which depends on extensional viscosity. Similarly, the viscous and elastic components are beneficial for forming of blow moulded and foamed PP products.

35 The two major parameters that contribute to the viscoelastic behaviour are molar mass and molar mass distribution.

Melt strength means the force needed in extension to break the polymer melt. Linear polymers generally have considerably low melt strength. For polypropylene the most determining parameter is the molar mass: the higher the molecular mass of the material, the higher the melt strength.

5

Known in the art (WO 94/26794, PoliBrasil) is a method of preparing high melt strength polypropylene by a continuous process, wherein propylene is polymerized in at least two polymerization vessels by adding discrete amounts of hydrogen as a molar mass regulator to at least one of the polymerization vessels to produce high melt strength polypropylene.

10

The resulting polymer product contains 10 to 35 % of a higher molar mass portion and 65 to 90 % of a lower molar mass portion. The process equipment comprises five slurry reactors connected in series, the amount of hydrogen fed to the polymerization vessels being successively reduced from the second polymerization vessel on.

15

In rheological terms, the polypropylene of WO 94/26794 is unsatisfactory as a high melt strength resin, because the zero viscosity and the shear thinning behaviour are rather similar to that of a conventional polypropylene used for comparison in the Examples of the reference. Thus, the products are not well suited to the manufacture of, e.g., foamed articles. Very little information on the catalysts used is given; in particular there is no

20

teaching on hydrogen sensitivity of the catalyst or on any catalyst donors employed. Further, because of the afore-mentioned process configuration, no hydrogen can be removed between the reactors, which limits the possibilities of varying the process conditions of the polymerization reactors. Thus, e.g., the high molar mass portion has to

25

be made in the first reactor and the low molar mass portion in the second reactor. In a slurry process (such as the one of WO 94/26794), carried out e.g. in heptane slurry, generally a TiCl_3 catalyst having low activity is used, because the activity level has to be controlled in the process due to limited heat transfer in these kind of processes.

30

Another patent, EP 0 498 603 (Amoco), describes a multireactor process for preparing polypropylene having a broad molecular mass distribution which, expressed as the ratio M_w/M_n , lies in the range of 7 to 11. This high stiffness polypropylene, which has high shear thinning, is suitable for, e.g. moulding thin walled products. The manufacture of foamed articles is not mentioned and the propylene polymers of the reference do not

35

exhibit the necessary melt strength for the production of light-weight polymer foams with closed cells.

Still another prior art publication, EP 0 573 862 (Himont), describes a polypropylene of very broad molecular mass distribution, $M_w/M_n > 20$. The polypropylene is claimed to have improved mechanical properties of high stiffness and improved resistance due to the broad MWD. The prior art process proposed for preparing the product comprises a multi-reactor gas phase process. The products are suggested for extrusion in thin sheets to be subjected to thermoforming. Once again, there is no suggestion of any use in foamed articles with closed cells and the polymers would not appear to have sufficiently high melt strength.

Summary of the Invention

It is an object of the invention to eliminate the problems related to the prior art and to provide novel linear homopolymers and copolymers of polypropylene having high melt strength and suited for, in particular, the preparation of foamed, blow moulded or thermoformed articles.

A second object of the invention is to provide a novel process for preparing high melt strength linear polypropylene without post-polymerization of the polymer.

It is still a further object of the present invention to provide extruded and moulded products comprising high melt strength polypropylene.

These and other objects, together with the advantages thereof over known processes and products, which shall become apparent from the specification which follows, are accomplished by the invention as hereinafter described and claimed.

The invention is based on tailoring the molar mass distribution of the polymer in a two-step polymerization process by using different amounts of a molar mass regulator in at least two of the polymerization steps. In connection with the present invention it has been found that particularly useful products, suitable for, e.g. blow moulding, thermoforming and foaming, can be obtained by using Ziegler-Natta catalysts or metallocene catalysts having decreased chain transfer sensitivity. The hydrogen sensitivity of the catalysts and its impact on the melt strength of the polymers are not discussed in any of the above-mentioned references.

Basically, the present invention comprises the steps of

- subjecting propylene and optionally other olefins to polymerization in a plurality of polymerization reactors connected in series,
- using different amounts of hydrogen and possibly comonomers as molar mass modifiers or regulators in at least two of the reactors, and
- carrying out the polymerization reaction in the presence of a catalyst system of low chain transfer sensitivity.

In connection with the present invention, it has been found that the high molar mass portion of the polymer product should be so high that a high viscosity at low shear stress rates approaching zero can be obtained. When the melt flow ratio, MFR_2 , of the product is below approx 3 g/10 min, a nominal viscosity at low shear rate (< 0.1 1/s) of about 40,000 Pa*s or more will provide a product having excellent melt strength which can be used for manufacturing foams and similar articles.

Thus, a polypropylene homopolymer or copolymer according to the present invention has improved melt strength, observed also in dynamic rheology as increased nominal viscosity at low shear rates of less than 0.1/sec. Its molar mass distribution (M_w/M_n) is > 6 and MFR_2 is 0.1 to 20, in particular below 5 and a nominal viscosity of more than 40,000 Pa*s.

More specifically, the polymer according to the invention is characterized by what is stated in the characterizing part of claim 1.

For obtaining a product of the above kind the catalyst system used should be capable of catalyzing the formation of high molar mass polymerization products (homo- or copolymers of propylene) having a MFR_2 of less than 0.1 g/10 min, in particular less than 0.04 g/10 min. It should also be capable of catalyzing the formation of low or medium molar mass polymerization products having a MFR_2 of 0.5 g/10 min or more.

The preferred catalyst system used comprises a Ziegler-Natta catalyst having a solid catalyst component, a cocatalyst component and an external donor. The chain transfer sensitivity of such a Ziegler-Natta catalyst can be reduced by using an external donor which is strongly coordinating. These kinds of donors are in particular represented by the general formula I



wherein R' represents a branched or cyclic aliphatic, or aromatic group.

Another preferred catalyst system comprises metallocene catalysts.

- 5 More specifically, the process according to the present invention is mainly characterized by what is stated in the characterizing part of claim 8.

10 The present homo- or copolymers can be used for the manufacture of polymer foams, thermoformed products, films and sheets, products prepared by blow molding or stretch blow molding, extrusion coating and injection moulding.

15 The invention achieves a number of considerable advantages which will appear from the following description. However, it should be particularly pointed out that the present linear high melt strength PP exhibits excellent forming stability in melt phase because of the very high molar mass fraction resulting in high viscosity at low shear rates.

Good extrusion behaviour is obtained due to a broad molar mass distribution, which results in a lowered viscosity at high shear rates.

20 In extrusion coating, the present polymer will exhibit reduced melt resonance at high draw ratios ($> 15 : 1$) and reduced neck-in. Products made by pipe extrusion and moulding will have improved properties, such as enhanced creep resistance. The polymer will also possess improved sag resistance, which is important in the thermoforming process. Furthermore, the material can be used to provide fibres of high resilience. In the
25 calendering process, the present material will exhibit less adhesion to the heat rolls and improved grain retention.

Next, the invention will be more closely examined with the aid of the following detailed description and with reference to the attached drawings.

30

Brief Description of the Drawings

Figure 1 depicts in a schematic fashion the process configuration of a first preferred embodiment of the invention;

35 Figure 2 depicts the process configuration of a second preferred embodiment of the invention;

Figure 3 shows a haul-off equipment which can be used for measuring melt strength; and Figure 4 shows the shear thinning behaviour at +200 °C for three polypropylene products according to the present invention compared with a commercial unimodal polypropylene.

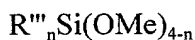
5 Detailed Description of the Invention

Definitions

10 The expression "using different amounts of hydrogen as a molar mass modifier" means in practice that only small amounts, if any, of hydrogen are employed in at least one reactor, whereas significant amounts, typically up to 10 to 15 mol hydrogen per kmol of propylene are used in at least one of the other reactors. As a result, a product containing a high molar mass portion and a low molar mass portion as obtained.

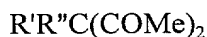
15 By "low (or reduced) chain transfer sensitivity" is meant that the number of chain transfer reactions occurring in the presence of the present catalyst system is smaller than the corresponding amount of chain transfer reactions in the presence of conventional catalyst with external electron donors, such as donor C, cyclohexyl methyl dimethoxy silane, and others.

20 "Strongly coordinating donor" designates donors which form relatively strong complexes with catalyst surfaces, mainly with a MgCl_2 surface in the presence of aluminium alkyl and TiCl_4 . The donor components are characterized by strong complexation affinity towards catalyst surfaces and a sterically large and protective hydrocarbon (R'). Strong coordination with MgCl_2 requires oxygen-oxygen distance of 2.5 to 2.9 Å [Albizzati et al., Macromol. Symp. 89 (1995) 73-89]. Typically this kind of donors has the structure of the general formula II



30 wherein R''' is a branched aliphatic or cyclic or aromatic group, and n is 1 or 2, preferably 2. [Härkönen et al., Macromol. Chem. 192 81991) 2857-2863].

Another group of such donors are 1,3-diethers having the formula III



wherein R' and R" are the same or different and stand for a branched aliphatic or cyclic or aromatic group.

By "Melt Flow Rate" or abbreviated "MFR" is meant the weight of a polymer extruded through a standard cylindrical die at a standard temperature in a laboratory rheometer carrying a standard piston and load. MFR is a measure of the melt viscosity of a polymer and hence also of its molar mass. The abbreviation "MFR" is generally provided with a numerical subindex indicating the load of the piston in the test. Thus, e.g., MFR₂ designates a 2.16 kg load and MFR₁₀ a load of 10.0 kg. MFR can be determined using, e.g., by one of the following tests: ISO 1133 C4, ASTM D 1238 and DIN 53735.

The overall process

The present invention concerns a multistage process for homo- or copolymerizing propylene to provide a product having high melt strength and a bimodal molar mass distribution comprising a high molar mass portion and a low or medium molar mass portion.

Propylene is polymerized in the presence of a Ziegler-Natta catalyst at an elevated temperature. Polymerization is carried out in a series of polymerization reactors selected from the group of slurry and gas phase reactors. Bulk polymerization in loop reactors is a particularly preferred embodiment of polymerization in slurry reactors. The high molar mass portion and the low or medium molar mass portion of the product can be prepared in any order in the reactors. If desired, the molar mass of the polymer can be raised by feeding a comonomer, such as ethylene, into the reactor.

The production split between the high molar mass polymerization reactor and the low or medium molar mass polymerization reactor is 5 - 95 : 95 - 5. Preferably, 50 to 90 %, in particular, 70 to 90 %, of the propylene homopolymer or copolymer is produced having a MFR₂ of 0.5 g/10 min or more and to provide a low or medium molar mass portion and 50 to 5 %, in particular 30 to 10 %, of the propylene homopolymer or copolymer is produced to provide a high molar mass portion having a MFR₂ of less than 0.1 g/10 min, in particular less than 0.04 g/10 min.

After preparation of the bimodal propylene homo- or copolymer, the properties of the

product can be modified by copolymerizing propylene with comonomers that will give the product properties of improved impact strength, for example. This modification can be carried out in polymerization reactors connected in series with the actual polymerization reactors.

5

The high melt strength polymer according to the invention comprises 50 to 95 % of a low or medium molar mass portion and 5 to 50 of a high molar mass portion and the Melt Flow Rate (MFR₂) of the homopolymer is about 0.1 to 20 g/10 min, preferably less than 10 g/10 min and in particular less than 5 g/10 min. The nominal viscosity of the polymer product is more than 40,000 Pa*s, in particular more than 45,000 Pa*s, at a Melt Flow Rate of 0.1 to 2 g/10 min. The melt strength of the polymer, determined by the haul-off method (as the highest value with highest possible haul-off speed before the polymer string is broken) is generally over 2.5, preferably 3 g or higher, in particular at least 5 g.

10

The main features of the present invention are the catalyst used, the polymerization reactor system and the properties of the polymer. These will be discussed in the following.

15

The catalyst

20

25

As catalyst any stereospecific catalyst for propylene polymerization can be used, which will yield high molecular mass at low hydrogen-to-propylene ratios and low or medium molecular mass at high hydrogen-to-propylene ratios at the typical conditions prevailing in polymerization reactors. Thus, the catalyst should preferably be capable of catalyzing the formation of the high molar mass product and the low or medium molar mass product at a pressure of 10 to 100 bar, in particular 25 to 80 bar, and at a temperature of 40 to 110 °C, in particular 60 to 100 °C. The catalyst has decreased sensitivity to chain transfer reactions and preferably contains an external, strongly coordinating donor.

30

Generally, the catalyst comprises a catalyst component, a cocatalyst component, an external donor, the catalyst component of the catalyst system primarily containing magnesium, titanium, halogen and an internal donor.

Examples of suitable catalyst systems are described in, for example, Finnish Patents Nos. 86866, 96615 and 88047 and 88048.

35

One particularly preferable catalyst, which can be used in the present invention, is

disclosed in FI Patent No. 88047. Another preferred catalyst is disclosed in Finnish Patent Application No. 963707.

A catalyst system useful in the present process can be prepared by reacting a magnesium halide compound with titanium tetrachloride and an internal donor. The magnesium halide compound is, for example, selected from the group of magnesium chloride, a complex of magnesium chloride with a lower alkanol and other derivatives of magnesium chloride. $MgCl_2$ can be used as such or it can be combined with silica, e.g. by absorbing the silica with a solution or slurry containing $MgCl_2$. The lower alkanol used can be preferably methanol or ethanol, particularly ethanol.

The titanium compound used in the preparation of the procatalyst is preferably an organic or inorganic titanium compound, having an oxidation state of titanium of 3 or 4. Also other transition metal compounds, such as vanadium, zirconium, chromium, molybdenum and tungsten compounds can be mixed with the titanium compound. The titanium compound usually is halide or oxyhalide, an organic metal halide, or a purely metal organic compound, in which only organic ligands have been attached to the transition metal. Particularly preferable are the titanium halides, especially $TiCl_4$. Preferably the titanation is carried out in two or three steps.

The alkoxy group of the phthalic acid ester used as an internal donor comprises at least five carbon atoms, preferably at least 8 carbon atoms. Thus, as the ester can be used for example propylhexyl phthalate, dioctyl phthalate, dinonyl phthalate, diisodecyl phthalate, di-undecyl phthalate, ditridecyl phthalate or ditetradecyl phthalate. The molar ratio of phthalic acid ester and magnesium halide is preferably about 0.2.

The transesterification can be carried out e.g. by selecting a phthalic acid ester - a lower alcohol pair, which spontaneously or with the aid of a catalyst, which does not damage the procatalyst composition, transesterifies the catalyst at an elevated temperatures. It is preferable to carry out the transesterification at a temperature, which lies in the range of 110 to 150 °C, preferably 120 to 140 °C.

The catalyst prepared by the method above is used together with an organometallic cocatalyst and with an external donor. Generally, the external donor has the formula IV



wherein

R and R' can be the same or different and they stand for a linear, branched or cyclic aliphatic, or aromatic group;

5 R" is methyl or ethyl;

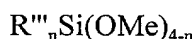
n is an integer 0 to 3;

m is an integer 0 to 3; and

n+m is 1 to 3.

10 The aliphatic groups in the meanings of R and R' can be saturated or unsaturated. Linear C₁ to C₁₂ hydrocarbons include methyl, ethyl, propyl, butyl, octyl and decanyl. As examples of suitable saturated branched C₁₋₈ alkyl groups, the following can be mentioned: isopropyl, isobutyl, isopentyl, tert-butyl, tert-amyl and neopentyl. Cyclic aliphatic groups containing 4 to 8 carbon atoms comprise, e.g., cyclopentyl, cyclohexyl,
15 methyl cyclopentyl and cycloheptyl.

According to the invention, particularly preferred donor are characterized as strongly coordinating donors which form relatively strong complexes with catalyst surface, mainly with MgCl₂ surface in the presence of aluminium alkyl and TiCl₄. The donor components
20 are characterised by a strong complexation affinity towards catalyst surface and a sterically large and protective hydrocarbon. Typically a donor of this kind has the structure V



25 wherein R''' is branched aliphatic or cyclic or aromatic group, and n is 1 or 2, preferably 2. [Makromol Chem. 192 (1991) p. 2857-2863, Härkönen et.al.]

In particular, the external donor is selected from the group consisting of dicyclopentyl dimethoxysilane, diisopropyl dimethoxysilane, di-isobutyl dimethoxysilane, and di-t-butyl dimethoxysilane.
30

An organoaluminum compound is used as a cocatalyst. The organoaluminium compound is preferably selected from the group consisting of trialkylaluminium, dialkyl aluminium chloride and alkyl aluminium sesquichloride.

35 Optionally any metallocene catalyst of capable of catalyzing the high and low or medium

molar mass product can be used. The hydrogen sensitivity of the metallocene catalyst can be different from that of a Ziegler-Natta catalyst which means that different hydrogen concentrations can be employed in the reactors.

5 The metallocene catalyst comprises a metallocene/activator reaction product impregnated in a porous support at maximum internal pore volume. The catalyst complex comprises a ligand which is typically bridged, and a transition metal of group IVA...VIA. The catalytic metal is typically halide, and aluminium alkyl. The ligands can belong to group of heterocyclic substituted or unsubstituted compounds, e.g. indocenes, naftenes, or any other
10 bulky compound which can control the stereoselectivity of the catalyst especially when ligands are bridged together with silane or other chemical bond. The activator is selected from a group in which are derivatives of water and aluminium alkyls e.g. trimethyl aluminium, triethyl aluminium, and tri t-butyl aluminium, or another compound capable of activating the complex. The metallocene/activator reaction product, a solvent capable of
15 dissolving it, and a porous support are brought into mutual contact, the solvent is removed and the porous support is impregnated with the metallocene/activator reaction product, the maximum amount of which corresponds to the pore volume support, cf. International PCT Application No. PCT/FI94/00499.

20 One typical structure of metallocene compound having decreased sensitivity to chain transfer reactions is bridged bis(2-R-4-R'-indenyl) M Cl₂, wherein both R and R' are aliphatic, cycloaliphatic or aromatic hydrocarbons having 1 to 18 C atoms, R' is typically phenyl or naphthyl, and R is typically methyl or ethyl. M is a transition metal, typically titanium, zirconium or hafnium. R and R' may contain heteroatoms, such as silicon,
25 nitrogen, phosphorous or germanium. The bridge between the indenyls is made from 1 to 3 atoms, such as carbon, silicon, nitrogen, phosphorous or germanium. A typical bridge is dimethylsilyl or ethyl. An example of such a metallocene compound is dimethylsilyl-bis(2-methyl-4-phenyl-indenyl)zirconium dichloride.

30 **Prepolymerization**

The catalyst can be prepolymerized prior to feeding into the polymerization reactor. During prepolymerization the catalyst components are contacted for a short period with a monomer, such as an olefin monomer, before feeding into the reactor. Examples of suitable
35 systems are described in, e.g., Finnish Patent Application No. 961152. It is also possible, as described in Finnish Patent No. 95387, to carry out the prepolymerization in the

presence of a viscous substance, such as an olefinic wax or an oil/wax mixture, to provide a prepolymerized catalyst which is stable during storage and handling. The catalysts prepolymerized in wax will also allow for easy dosing of the catalyst into the polymerization reactors.

5

Polymerization process

The equipment of the polymerization process following the optional prepolymerization step can comprise a series of polymerization reactors of any suitable type producing propylene homo- or comonomers of different molar masses in the presence and absence, respectively, of hydrogen or another molar mass regulator. Thus, the polymerization reactor system can comprise a series of conventional stirred-tank slurry reactors, as described in WO 94/26794. Preferably the reactors used are selected from the group of loop and gas phase reactors and, in particular, the process employs at least one loop reactor and at least one gas phase reactor. It is also possible to use several reactors of each type, e.g. one loop reactor and two or three gas phase reactors, in series. In the following the reactor system will be described with particular reference to a system comprising one loop reactor (referred to as "the first reactor") and one gas phase reactor (referred to as "the second reactor"), in that order. However, it should be understood that the reactor system can comprise the reactors in any number and order.

In every polymerization step it is possible to use also comonomers selected from the group of ethylene, propylene, butene, pentene, hexene and alike as well as their mixtures. The use of comonomers other than ethylene is particularly preferred for the preparation of the low or medium molar mass portion.

In addition to the actual polymerization reactors used for producing the bimodal propylene homo- or copolymer, the polymerization reaction system can also include a number of additional reactors, such as pre- and/or postreactors. The prereactors include any reactor for prepolymerizing the catalyst and for modifying the olefinic feed, if necessary. The postreactors (also referred to as the "third", "fourth" etc. reactor) include reactors used for modifying and improving the properties of the polymer product, e.g. impact resistance, as described above. All reactors of the reactor system are preferably arranged in series.

According to the invention, the polymerization comprises the steps of

- subjecting propylene and optionally other olefins to a first polymerization or

- copolymerization reaction in a first reaction zone or reactor,
- recovering the first polymerization product from the first reaction zone,
 - feeding the first polymerization product to a second reaction zone or reactor,
 - feeding additional propylene to the second reaction zone,
 - 5 - subjecting the additional propylene to a second polymerization reaction in the presence of the first polymerization product to produce a second polymerization product, and
 - recovering the second polymerization product from the second reaction zone, and
 - separating and recovering the polypropylene from the second reaction product.

10 Thus, in the first step of the process, propylene with the optional comonomer(s) together with the catalyst is fed into the first polymerization reactor. Along with these components hydrogen as a molar mass regulator is fed into the reactor in the amount required for achieving the desired molar mass of the polymer. Alternatively, the feed of the first reactor

15 can consists of the reaction mixture from a previous reactor, if any, together with added fresh monomer, hydrogen, optional comonomer and cocatalyst. In the presence of the catalyst, propylene and the optional comonomer will polymerize and form a product in particulate form, i.e. polymer particles, which are suspended in the fluid circulated in the reactor.

20 The polymerization medium typically comprises the monomer (i.e. propylene) and/or a hydrocarbon, and the fluid is either liquid or gaseous. In the case of a slurry reactor, in particular a loop reactor, the fluid is liquid and the suspension of polymer is circulated continuously through the slurry reactor, whereby more suspension of polymer in particle

25 form in a hydrocarbon medium or monomer will be produced. According to a preferred embodiment, the first polymerization or copolymerization reaction is carried out in a reaction medium mainly consisting of propylene, Preferably at least 60 weight percent of the medium is formed by propylene.

30 The conditions of the slurry reactor are selected so that at least 10 wt-%, preferably at least 15 wt-%, of the whole production is polymerized in each slurry reactor. The temperature is in the range of 40 to 110 °C, preferably in the range of 70 to 100 °C. The reaction pressure is in the range of 25 to 100 bar, preferably 35 to 80 bar.

35 In slurry polymerization more than one reactor can be used in series. In such a case the polymer suspension in an inert hydrocarbon or in monomer produced in the slurry reactor

is fed without separation of inert components and monomers periodically or continuously to the following slurry reactor, which acts at lower pressure than the previous slurry reactor.

- 5 The polymerization heat is removed by cooling the reactor by a cooling jacket. The residence time in the slurry reactor must be at least 10 minutes, preferably 20-100 min for obtaining a sufficient degree of polymerization.

10 According to one embodiment of the invention light inert hydrocarbons are fed to the reactor. Examples of such hydrocarbons are iso-butane, n-butane and isopentane. The light, inert hydrocarbon in the polymerization mixture lowers the pressure required in the reactor. The increased catalyst activity at relatively high temperature compensates for the decreased activity due to lowered concentration of propylene.

- 15 As discussed above, if a low or medium molar mass polypropylene is the desired product, hydrogen is fed into the reactor. Hydrogen can be added to the reactor at ratio of at least 1.5 mol H₂/kmol propylene, preferably 1.5-15 mol H₂/kmol propylene.

20 If the first polymerization reaction is carried out in the presence of hydrogen, then the pressure of the first polymerization product including the reaction medium is preferably reduced after the first reaction zone in order to evaporate volatile components of the product, e.g. in a flash tank. An overhead stream obtained from the evaporation of volatile components is condensed, and a concentrated fraction of propylene is recovered from the condensed stream and recirculated to the first reactor. As a result of the
25 flashing, the product stream containing the polypropylene is freed from hydrogen and can be subjected to a second polymerization in the presence of additional propylene to produce a high molar mass polymer. If, again, the high molecular mass polymer is produced in the first reactor, then the polymerization product can be directly conducted as such to the following reactor.

30 The second reactor is preferably a gas phase reactor, wherein propylene and optionally comonomers are polymerized in a gaseous reaction medium.

35 The gas phase reactor can be an ordinary fluidized bed reactor, although other types of gas phase reactors can be used. In a fluidized bed reactor, the bed consists of the formed and growing polymer particles as well as still active catalyst come along with the polymer

fraction. The bed is kept in a fluidized state by introducing gaseous components, for instance monomer on such flowing rate which will make the particles act as a fluid. The fluidizing gas can contain also inert carrier gases, like nitrogen and also hydrogen as a modifier. The fluidized gas phase reactor can be equipped with a mechanical mixer.

5

The gas phase reactor used can be operated in the temperature range of 50 to 115 °C, preferably between 60 and 110°C and the reaction pressure between 10 and 40 bar and the partial pressure of monomer between 2 and 30 bar.

10

The pressure of the second polymerization product including the gaseous reaction medium can then be released after the second reactor in order optionally to separate part of the gaseous and possible volatile components of the product, e.g. in a flash tank. The overhead stream or part of it is recirculated to the second reactor.

15

If desired, the polymerization product can be fed into a second gas phase reactor in which a rubbery copolymer is provided by a third (co)polymerization reaction to produce a modified polymerization product. This third polymerization reaction will give the polymerization product properties of e.g. improved impact strength. The step of providing an elastomer can be performed in various ways. Thus, preferably an elastomer is produced by copolymerizing at least propylene and ethylene into an elastomer. The conditions for the copolymerization are within the limits of conventional EPM production conditions such as they are disclosed, e.g., in Encyclopedia of Polymer Science and Engineering, Second Edition, Vol. 6, p.545-558. A rubbery product is formed if the ethylene repeating unit content in the polymer lies within a certain range. Thus, preferably, ethylene and propylene are copolymerized into an elastomer in such a ratio that the copolymer contains from 10 to 70 % by weight of ethylene units. In particular, the ethylene unit content is from 30 to 50 % by weight of the copolymer propylene/ethylene elastomer. In other words, ethylene and propylene are copolymerized into an elastomer in a molar ratio of ethylene-to-propylene of 30/70 to 50/50.

20

25

30

The elastomer can also be provided by adding a ready-made or natural elastomer to the polymer product of the first gas phase reactor.

35

Summarizing what has been stated above, one particularly preferred embodiments of the invention comprises the following steps:

- propylene is polymerized in a loop reactor at a pressure of 25 to 80 bar, at a

temperature of 60 to 100 °C to provide a low or medium molar mass polymerization product,

- the polymerization product of the loop reactor is recovered and conducted to a flash tank, wherein an overhead product containing hydrogen and non-reacted propylene is separated from a bottom product containing polymerized solids,
- at least a part of the overhead product is recycled to the loop reactor,
- the bottom product is conducted to a gas phase reactor,
- additional propylene and, optionally ethylene, is fed to the gas phase reactor,
- the additional propylene is subjected to polymerization at a pressure of 20 bar or more to provide a high molar mass polymerization product,
- the polymerization product of the gas phase reactor is recovered and conducted to a flash tank, wherein the pressure of the product is reduced to produce an overhead product containing hydrogen and non-reacted propylene and a bottom product primarily containing polymerized solids, and
- polypropylene polymer is recovered from a part of the bottom product of the flash tank.

According to the present, at least a part of the overhead product can be recycled to the loop reactor, but it is also possible to conduct the overhead product directly to the gas phase reactor after at least a partial separation of hydrogen.

According to the second particularly preferred embodiment:

- propylene and optional ethylene is polymerized in a loop reactor at a pressure of 25 to 80 bar, at a temperature of 60 to 100 °C to provide a high molar mass polymerization product,
- the polymerization product and reaction medium of the loop reactor is conducted directly to a gas phase reactor fluid bed,
- additional propylene, if desired, is fed to the gas phase reactor,
- propylene is subjected to polymerization in the gas phase reactor at a pressure of 20 bar or more to provide a low or medium molar mass polymerization product,
- the polymerization product of the gas phase reactor is recovered and conducted to a flash tank, wherein the pressure of the product is reduced to produce an overhead product containing hydrogen and non-reacted propylene and a bottom product primarily containing polymerized solids,
- at least a part of the overhead product is recycled to the gas phase reactor, and
- polypropylene polymer is recovered from a part of the bottom product of the flash

tank.

The optional addition of ethylene in the reactor producing a high molecular mass fraction increases the actual molar mass produced in the reactor. Ethylene can also be added to the reactor producing a low MW fraction and the increased MW can be compensated with extra hydrogen.

These two preferred embodiments are also depicted in the attached drawings, which illustrate the particular configuration of process equipment used. The reference numerals used in the drawings are common for both embodiments, since the apparatus is largely the same. The numerals refer to the following apparatuses:

- 1 prepolymerization reactor
- 30 catalyst reservoir
- 31 feeding device
- 32 diluent
- 33 catalyst/diluent mixture
- 34 monomer
- 35 cocatalyst and possible donors
- 40 loop reactor
- 42 diluent feed (optional)
- 43 monomer feed
- 44 hydrogen feed
- 45 comonomer feed (optional)
- 46 back to the loop reactor 40 through the line 46
- 47 one or several exhaust valve
- 48 product transfer line
- 50 flash separator
- 51 recovery unit and off gas vent
- 52 removing line
- 60 gas phase reactor
- 61 gas transfer line
- 62 compressor
- 63 monomer feed
- 64 comonomer feed
- 65 hydrogen feed
- 66 transfer line
- 67 product transfer line
- 68 recovery system

Turning to Figure 1, it can be noted that catalyst from reservoir 30 is fed to the feeding device 31 together with diluent from line 32. The feeding device 31 feeds the catalyst/diluent mixture into the prepolymerization chamber 1 via line 33. Monomer is fed through 34 and cocatalyst and possible donors can be fed into the reactor 1 through

From the prepolymerization chamber 1 the prepolymerized catalyst is removed preferably directly through line 36 and transferred to a loop reactor 40. In the loop reactor 40 the polymerization is continued by adding a diluent from the line 42, monomer from line 43, hydrogen from line 44 and an optional comonomer from line 45 through the line 45 through the line 46. An optional cocatalyst can also be introduced into the loop reactor 40.

In the lower part of the gas phase reactor 60 there is a bed consisting of polymer particles, which will be kept in a fluidized state in an ordinary way by circulating the gases removed from the top of the reactor 60 through line 61, compressor 62 and a heat exchanger (not presented) to the lower part of the reactor 60 in an ordinary way. The reactor 60 is advantageously, but not necessarily, equipped with a mixer (not presented). To the lower part of the reactor 60 can be led in a well known way monomers from line 63, optionally comonomer from line 64 and hydrogen from the line 65. The product will be removed from the reactor 60 continually or periodically through the transfer line 66 to the recovery system 68. The overhead product of the recovery system is recirculated to the gas reactor.

The product

By means of the present invention, a propylene homopolymer or copolymer is produced having improved shear thinning and melt elasticity values. The higher the shear thinning index, the higher the melt elasticity. The properties are manifested in dynamic rheology as increased nominal viscosity at shear rates $< 0.1/\text{sec}$. It is obvious that both properties are influenced by high molar mass tails. The high molar mass tails also give improve melt strength characterized by the haul-off method. The molar mass distribution of the polymer is > 6 , preferably > 8 , and the Melt Flow Rate (MFR_2) 0.1 to 20 g/10 min. The melt strength of the polymer product (determined by the haul-off method described below) is over about 2 to 2.5 g, preferably it is at least 3 g and, in particular it is over 5 g. By feeding (e.g. about 0.01 to 5 mole-%) ethylene as a comonomer to the reactor wherein the high molar mass is produced, it is possible further to increase the melt strength.

It should be noticed that due to the specific composition of the present bimodal polymers, comprising a high molar mass portion with a rather high molar mass, the melt strength of the present polymers is about 1.5 times (or more) higher than for corresponding unimodal polymers having an equal or similar Melt Flow Rate.

MFR measurements show that polymerization of propylene without any addition of hydrogen into the reactor vessel results in the formation of polypropylenes with very high molar mass. In addition, the molar mass is dependent on the catalyst system used as well as on the polymerization conditions.

Typically, the MFR_2 of polypropylene made in liquid monomer essentially or entirely in the absence of hydrogen using a catalyst system containing a strongly complexing donor is less than 0.02 g/10 min, e.g. approximately 0.01 g/10 min, whereas a corresponding catalyst system with a less strongly complexing donor, under the same conditions, typically gives polypropylene with a MFR_2 of > 0.04 .

Molar mass distributions are broad, $M_w/M_n > 6$, preferably > 8 . It should be noted that when the portion of the high molar mass is small and the molar mass very high, the MWD can be difficult to determine using standard methods such as GPC.

The homopolymer or copolymer described above can be used for the manufacture of

polymer foams, thermoformed products, films and sheets, and products prepared by blow molding or stretch blow molding, extrusion coating or injection moulding.

In particular, the novel propylene polymers can be employed for manufacturing foamed materials, such as beads, strands, sheets, laminated sheets, netting and profiles. The present polymers will provide foams with closed cells and densities in the range from 20 to 400 kg/m³, in particular about 20 to 60 kg/m³. The foamed materials can be used, for example, as moulded parts in the automotive industry, in flooring, for pipe insulation, or in sheets for thermoforming.

The novel polymers can be processed into foamed materials by methods known *per se*. Thus, they can be extrusion foamed in a extruder, such as a single screw extruder, twin screw extruder or a tandem extruder. The polymers according to the invention can also be foamed by using a physical blowing agent or a chemical blowing agent or a combination thereof. As examples of physical blowing agents the following can be mentioned: CO₂, different hydrocarbons (including propane and butane), H₂O and CFC's and mixtures thereof. The chemical blowing agent can be selected from the group consisting of inorganic carbonates, citric acid and azodicarbon amide and mixtures thereof. A bubble initiator, such as talc or citric acid, can be used to improve the foam structure.

The present polymers and copolymers of propylene can be blended and optionally compounded with additives and adjuvants conventionally used in the art. Thus, suitable additives include antistatic agents, flame retardants, light and heat stabilizers, pigments and carbon black. Fillers such as mica can also be used. Further, the polymers and copolymers can be blended with other polymers, in particular other polyolefins, such as LD-, LLD-, MD- and HD-polyethylenes and polybutylene.

Examples

The following non-limiting examples illustrate the invention.

In the Examples, the Melt Flow Rate was measured at 230 °C using piston having loads of 2.16 kg and 10 kg. The abbreviations MFR₂ and MFR₁₀ are used for designating the melt flow rate measured at these conditions.

Dynamic rheological measurements were performed by Rheometrics System IV rheometer

in a frequency range from 0.01 to 400 Hz at 200 °C.

Shear thinning index (SHI) is defined as a ratio of shear viscosities measured at two different shear stresses: $SHI = \eta(G^* = 2 \text{ kPa})/\eta(G^* = 50 \text{ kPa})$. η_0 stands for the viscosity measured at zero shear stress.

Melt strength has been determined by rheologic measurements on a "haul-off equipment". (see Figure 3). The haul-off equipment comprises a rosand capillary 101, a rheometer system 102, a balance 103, a pulley block combination 104 and a haul-off unit 105. A polymer string 106 formed from polymer melt in the capillary 101 is suspended from the rheometer system 102 and extended by the haul-off unit 105. The balance 103 will register the extensional viscosity (or melt strength) in terms of weight (g). Melt strength is determined as the highest value with highest possible haul-off speed before the polymer string is broken.

Example 1

Polymerization of propylene on laboratory scale

Propylene was polymerized on bench scale by producing a polymer fraction of high molar mass in a first polymerization step without any addition of hydrogen and in a second polymerization step, which was carried out in the presence of hydrogen in order to broaden the molar mass distribution and increase the final MFR to required level.

The polymerizations were carried out in the presence of a Ziegler-Natta catalyst with an external electron donor corresponding to the general formula I and V above (dicyclopentyl dimethoxy silane, in the following abbreviated "D"). The catalyst was prepared according to the method described in FI Patent No. 88047. A polymer prepared in the presence of a corresponding catalyst system containing a conventional donor corresponding to the general formula IV (cyclohexyl methyl dimethoxy silane or abbreviated "C") was tested. As a reference, a commercial polypropylene was used. Polymerizations were made in liquid propylene where the first polymerization step was made without any hydrogen in the feed and the second step in the presence of hydrogen.

The results of the polymerizations are summarized in Table 1.

Table 1. Results of the two step polymerizations

Property	A	B	C	Normal commercial PP
Catalyst donor	D	D	C	C
MFR ₂ , g/10 min	1.0	1.8	1.2	1.9
H ₂ (1st step)/ H ₂ (2nd step) / mmol	0/500	0/800	0/96	-
MFR ₂ (1st step) MFR ₂ (2nd step)	0.008/24	0.008/58	0.04/14	-
Split min/min	30/70 30/15	30/70 30/17	30/70 30/21	100
Activity, kg/g _{cat}	28.5	27.3	20.9	-
SHI (2/50)	180	240	24	5.9
η (2 kPa) (Pa*s)	170,000	48,000	46,000	10,000
Melt strength, g	6	10	3	1.4

Polymerization conditions: Al/Ti = 250; Al/D = 10; propylene = 840 g; T_p = 70 °C

As will appear from the above data, the polymerization of propylene without any addition of hydrogen into the reactor vessel results in the formation of polypropylenes with very high molar mass.

The viscosity curves of the above four different polypropylene polymers are also depicted in Figure 4.

As will appear from Figure 4, the viscosity of the polymer with decreasing shear rate was most pronounced with the polymers of the present invention. Thus, SHI of the present polymers were 180 (MFR₂ = 1.0 g/10 min) and 240 (MFR₂ = 1.8 g/10 min), which are much higher than that of standard grade with narrow MWD (SHI = 5.9, MFR₂ = 1.9 g/10 min). The polymer C prepared using a less strongly coordinating donor showed much lower shear thinning with increasing shear rate (SHI = 24, MFR₂ = 1.2 g/10 min). The different behaviour of the tested catalysts indicates that the molar mass distribution of the preferred polymers is much broader than the MWD of the polymer made with a less strongly coordinating donor.

As a result of the fraction of very high M_w , the present polymers have a much higher melt strength than reference grade polymers. Thus, the melt strength of polymer B, measured by the haul-off method, was 10, whereas the reference polymer D had a melt strength of 1.4. The higher melt strength of the present polymers is due to the broader MWD of the polymer and the presence of a very high molar mass fraction.

Example 2

Two-stage continuous polymerization of propylene:

Propylene homopolymers were produced in a pilot plant having a loop reactor and an agitated fluid bed gas-phase reactor connected in series. The catalyst, cocatalyst, and an external donor were fed into the loop reactor. The reaction medium, including hydrogen, of the loop was flashed away before the solid polymer containing the active catalyst entered the gas-phase reactor.

A prepolymerized $MgCl_2$ supported Ti-catalyst (prepared according to FI Patent No. 88 047) was used in the polymerization. The cocatalyst was triethyl aluminium (TEA) and dicyclopentyl dimethoxy silane (DCPDMS) was used as an external donor. The Al/Ti mole ratio was 150 and Al/donor mole ratio 5.

In the first stage (loop reactor), a low molar mass propene homopolymer was produced and polymerization was continued in the second stage (gas-phase reactor) which produced high molar mass propene homopolymer. Hydrogen feed was used in the loop reactor to control the MFR of the product. No additional hydrogen was fed to the gas-phase reactor. The MFR_2 of the loop product was 24 g/10 min. The Al/Ti mol ratio in the polymerization was 250 and Al/donor mol ratio 10. MFR_2 of the final product was 2.8 g/10 min. Production rate ratio between the first and the second reactor was 85/15. Polymerization temperature in the loop reactor was 70 °C and in the gas phase reactor 75 °C. The polymer had an η_0 of 55,300 Pa*s, a SHI (0/50) of 83 and a melt strength of 3 g.

Example 3

Two-stage continuous copolymerization of propylene and ethene

Polymerization of propylene in a loop reactor was carried out as in Example 2 but using higher hydrogen feed in order to adjust MFR_2 to 180 in the loop reactor. The reaction was continued in a second stage (a gas-phase reactor) where no hydrogen but additional

propylene and 2.5 mol-% ethylene were fed. The MFR_2 of the product was 2.6 g/10 min and ethylene content 0.8 wt-%. The production rate ratio between the first and the second reactor was 70/30. Polymerization temperatures and Al/Ti and Al/donor mol ratios were as in Example 2. The polymer had an η_0 of 116,300 Pa*s, a SHI (0/50) of 261 and a melt strength of 5 g.

Example 4

Two-stage continuous copolymerization of propylene and ethylene

Propylene copolymers were produced in pilot plant having a loop reactor and a fluid bed gas-phase reactor connected in series. The catalyst, cocatalyst and donor, as in Example 2, were fed into the loop reactor.

In the first stage (loop reactor), there was produced a high molar mass copolymer of propylene and ethylene having a MFR_2 of 0.03 g/10 min and ethylene content of 2.5 wt-%. Polymerization was continued in the second stage (gas-phase reactor) without flashing the non-reacted monomers between the reactors. In the gas-phase reactor the low molar mass homopolymer fraction was produced by adding additional propylene and hydrogen to the reactor. MFR_2 of the final product was 1.9 g/10 min and ethylene content 1.3 wt-%.

Production rate ratio between the reactors was 30/70. Polymerization temperatures and Al/Ti and Al/donor mol ratios were as in Example 2. The polymer had an η_0 of 199,400 Pa*s, a SHI (0/50) of 354 and a melt strength of 9 g.

Example 5

Two-stage continuous polymerization of propylene

Propylene homopolymers were produced in pilot plant having a loop reactor and a fluid bed gas-phase reactor connected in series. The catalyst, cocatalyst and donor, as in Example 2, were fed into the loop reactor.

In the first stage (loop reactor), there was produced a high molar mass homopolymer of propylene having a MFR_{10} of 0.2 g/10 min. Polymerization was continued in the second stage (gas-phase reactor) without flashing the non-reacted monomers between the reactors. In the gas-phase reactor the low molar mass homopolymer fraction was produced by feeding additional propylene and hydrogen to the reactor. The MFR_2 of the final product was 4.5 g/10 min. Production rate ratio between the reactors was 20/80. Polymerization

temperatures in the first and second step were 68 and 75 °C, respectively, the Al/Ti molar ratio was 250 and the Al/donor molar ratio was 20. The polymer had an η_0 of 49,000 Pa*s, a SHI (0/50) of 134 and a melt strength of 3 g.

5 Example 6

Two-stage continuous polymerization of propylene on bench scale

10 Propylene was polymerized on bench scale by producing a polymer fraction of high molar mass in a first polymerization step without any addition of hydrogen and in a second step which was carried out in the presence of hydrogen in order to broaden the molar mass distribution and increase the final MFR to required level.

15 The polymerization was carried out in the presence of a supported dimethylsilyl-bis(2-methyl-4-phenyl-indenyl)zirconium dichloride/methylaluminioxane catalyst.

Polymerization was made in liquid propylene at a temperature of 70 °C, the first 20 min polymerization step being carried out without any hydrogen in the feed and the second step (60 min) in the presence of 30 mmol hydrogen.

20 The activity of the catalyst was 0.81 kg/g cat and the MFR2 of the polymer was 1.2 g/10 min. The polymer had a molar mass distribution, M_w/M_n , of 10 and an excellent melt strength.

CLAIMS:

1. A linear polypropylene homopolymer or copolymer exhibiting a melt strength of at least 2.5 g, determined by the haul-off method, comprising a high molar mass portion and a low or medium molar mass portion, the high molar mass portion having a Melt Flow Rate (MFR₂) of less than 0.1 g/10 min and the low or medium molar mass portion having a Melt Flow Rate (MFR₂) of more than 0.5 g/10 min, the polymer having a Melt Flow Rate (MFR₂) of 0.1 to 20 g/10 min.

2. The polypropylene homopolymer or copolymer of claim 1 exhibiting improved melt strength observed in dynamic rheology as increased nominal viscosity at low shear rates of less than 0.1/sec, and having a MWD of more than 6.

3. The polypropylene homopolymer or copolymer of claim 1 or 2, exhibiting a melt strength of at least 5 g, determined by the haul-off method.

4. The polypropylene copolymer of any of claims 1 to 3, comprising 0.1 to 20 mole-% of ethylene residues.

5. Polymer product, comprising at least 50 % by weight of a polymer according to any of claims 1 to 4.

6. The polymer product according to claim 5, comprising 1 to 40 wt-% of a rubbery component.

7. The polymer product according to claim 6, wherein the rubbery component is selected from the group of elastomers produced by copolymerizing at least propylene and ethylene, ready-made elastomers and natural elastomers.

8. A process for preparing linear high melt strength propylene homopolymers and copolymers comprising the steps of

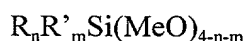
- subjecting propylene and optionally other olefins to polymerization in a plurality of polymerization reactors connected in series,
- employing different amounts of hydrogen as a molar mass modifier in at least two of the reactors, and
- carrying out the polymerization reaction in the presence of a catalyst system capable

of catalyzing the formation of a high molar mass polymerization product having a MFR₂ of less than 0.1 g/10 min and a low or medium molar mass polymerization product having a MFR₂ of more than 0.5 g/10 min.

9. The process according to claim 8, wherein the polymerization reaction is carried out in the presence of a catalyst system, which is capable of catalyzing the formation of a high molar mass polymerization product having a MFR₂ of less than 0.04 g/10 min and a low or medium molar mass polymerization product having a MFR₂ of more than 0.5 g/10 min.

10. The process according to any one of claim 8 or 9, wherein the catalyst is capable of catalyzing the formation of said high molar mass product and said low or medium molar mass product at a pressure of 25 to 80 bar and at a temperature of 60 to 100 °C in a loop reactor.

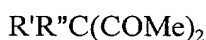
11. The process according to any one of claims 8 to 10, wherein the catalyst comprises a catalyst component, a cocatalyst component, an external donor and, optionally, an internal donor, the procatalyst component of the catalyst system containing magnesium, titanium, halogen and an electron donor, said catalyst system further containing an external donor having the general formula of



wherein R' and R can be the same or different and represent branched or cyclic aliphatic, or aromatic groups, and n and m are 0 or 1 and n + m is 1 or 2.

12. The process according to claim 11, wherein the external donor is selected from the group consisting of dicyclopentyldimethoxy silane, di-*tert*-butyldimethoxy silane, diisopropyldimethoxy silane and diisobutyldimethoxy silane.

13. The process according to any one of claims 9 to 10, wherein the catalyst comprises a catalyst component, a cocatalyst component, an external donor and, optionally, an internal donor, the procatalyst component of the catalyst system containing magnesium, titanium, halogen and an electron donor, said catalyst system further containing an external donor having the general formula of



wherein R' and R" are the same or different and stand for a branched aliphatic or cyclic or aromatic group.

5 14. The process according to claim 8 or 9, wherein the catalyst system comprises a metallocene catalyst.

10 15. The process according to claim 14, wherein the metallocene compound in the metallocene catalyst is bridged bis(2-R-4-R'-indenyl) M Cl₂, wherein both R and R' are aliphatic, cycloaliphatic or aromatic hydrocarbons having 1 to 18 C atoms, R' is typically phenyl or naphthyl, and R is a lower alkyl, M is a transition metal, and R and R' may contain heteroatoms, such as silicon, nitrogen, phosphorous or germanium, and the bridge between the indenyls comprises 1 to 3 atoms, such as carbon, silicon, nitrogen, phosphorous or germanium.

15 16. The process according to claim 15, wherein the metallocene compound is dimethylsilyl-bis(2-methyl-4-phenyl-indenyl)zirconiumdichloride.

20 17. The process according to any one of the preceding claims, wherein 5 to 50 % of the propylene homopolymer or copolymer has a MFR₂ < 0.1 g/10 min and 50 to 95 % of the propylene homopolymer or copolymer has a MFR₂ > 0.5 g/10 min.

25 18. The process according to any one of the preceding claims, wherein said reactors are selected from the group of loop reactors and gas phase reactors.

19. The process according to claim 18, wherein said reactors comprises at least one loop reactor and at least one gas phase reactor.

30 20. The process according to claim 19, wherein said reactors comprise a loop reactor and a gas phase reactor in that order.

35 21. The process according to any one of the preceding claims, wherein ethylene is fed into the reactor in which the higher molar mass component is produced in order to increase reactivity and molar mass of the component.

22. The process according to any one of the preceding claims, which comprises the steps of

- subjecting propylene and optionally other olefins to a first polymerization or copolymerization reaction in a first reaction zone,
- recovering the first polymerization product from the first reaction zone,
- feeding the first polymerization product to a second reaction zone,
- 5 – feeding additional propylene to the second reaction zone,
- subjecting the additional propylene to a second polymerization reaction in the presence of the first polymerization product to produce a second polymerization product, and
- recovering the second polymerization product from the second reaction zone, and
- 10 – separating and recovering the polypropylene from the second reaction product.

23. The process according to claim 22, wherein the first polymerization or copolymerization reaction is in a reaction medium, at least 60 weight percent of which is formed by propylene, producing a polymerization product in particulate form.

24. The process according to claim 22, wherein the first polymerization reaction is carried out in a loop reactor.

25. The process according to claim 23 or 24, wherein the pressure of the first polymerization product including the reaction medium is reduced after the first reaction zone in order to evaporate volatile components of the product.

26. The process according to claim 25, wherein the pressure of the first polymerization product is reduced in a flash tank.

27. The process according to claim 25 or 26, wherein an overhead stream obtained from the evaporation of volatile components is condensed, a concentrated fraction of propylene being recovered from the condensed stream and recirculated to the first reactor.

28. The process according to any one of the preceding claims, wherein the second reactor is a gas phase reactor, wherein propylene and optionally comonomers are polymerized in a gaseous reaction medium.

29. The process according to claim 28, wherein the pressure of the second polymerization product including the gaseous reaction medium is reduced after the second reactor in order to separate part of the gaseous and possible volatile components of the product.

30. The process according to claim 27, wherein the pressure of the second reactor is reduced in a flash tank.

31. The process according to claim 28, wherein the overhead stream or part of it is recirculated to the second reactor.

32. The process according to any one of claims 21 to 28, wherein the second polymerization product is fed into a third reactor and propylene is subjected to a third polymerization reaction to produce a third polymerization product.

33. The process according to claim 30, wherein the third polymerization reaction is carried out in a gas phase reactor in the presence of comonomers which give the third polymerization product properties of improved impact strength.

34. The process according to any one of claims 32 or 33, wherein the product of polymerization reaction is transferred to a fourth polymerization reaction zone.

35. The process according to claim 8, wherein

- propylene and optionally other olefins are polymerized in a loop reactor at a pressure of 25 to 80 bar, at a temperature of 60 to 100 °C to provide a low or medium molar mass polymerization product of $MFR_2 > 0.5$,
- the polymerization product of the loop reactor is recovered and conducted to a flash tank, wherein an overhead product containing hydrogen and non-reacted propylene is separated from a bottom product containing polymerized solids,
- the bottom product is conducted to a gas phase reactor,
- additional propylene and optionally other olefins are fed to the gas phase reactor,
- the additional propylene and optionally other olefins are subjected to polymerization at a pressure of 20 bar or more to provide a high molar mass polymerization product of $MFR_2 < 0.1$,
- the polymerization product of the gas phase reactor is recovered and conducted to a flash tank, wherein the pressure of the product is reduced to produce an overhead product containing hydrogen and non-reacted propylene and a bottom product primarily containing polymerized solids,
- at least a part of the overhead product is recycled to the gas phase reactor, and
- polypropylene polymer or copolymer is recovered from a part of the bottom product of the flash tank.

36. The process according to claim 35, wherein at least a part of the overhead product is recycled to the loop reactor.

37. The process according to claim 36, wherein the overhead product is conducted to the gas phase reactor after at least a partial separation of hydrogen.

38. The process according to claim 8, wherein

- propylene and optionally other olefins are polymerized in a loop reactor at a pressure of 25 to 80 bar, at a temperature of 40 to 100 °C to provide a high molar mass polymerization product of $MFR_2 < 0.1$,
- the polymerization product of the loop reactor is conducted directly to a gas phase reactor fluid bed,
- additional propylene and optionally other olefins are fed to the gas phase reactor,
- the additional propylene and optionally other olefins are subjected to polymerization at a pressure of 20 bar or more to provide a low or medium molar mass polymerization product of $MFR_2 > 0.5$,
- the polymerization product of the gas phase reactor is recovered and conducted to a flash tank, wherein the pressure of the product is reduced to produce an overhead product containing hydrogen and non-reacted propylene and a bottom product primarily containing polymerized solids,
- at least a part of the overhead product is recycled to the gas phase reactor, and
- polypropylene polymer or copolymer is recovered from a part of the bottom product of the flash tank.

39. The process according to any claims 8 to 13, wherein the catalyst system has been prepared by

- providing a procatalyst by reacting a magnesium halide compound, selected from the group consisting of magnesium chloride, a complex of magnesium chloride with ethanol and other derivatives of magnesium chloride, with titanium tetrachloride and an internal donor,
- providing as a cocatalyst an organoaluminium compound selected from the group consisting of trialkyl aluminium, dialkyl aluminium chloride and alkyl aluminium sesquichloride,
- providing an external donor having the general formula $R'_2(MeO)_2Si$, wherein R' stands for a cyclic or branched aliphatic, or aromatic group, for example dicyclopentyl dimethoxysilane or di-t-butyl dimethoxysilane, and, optionally,

- prepolymerizing a small amount of olefin by contacting the olefin with said procatalyst, cocatalyst and the external donor.

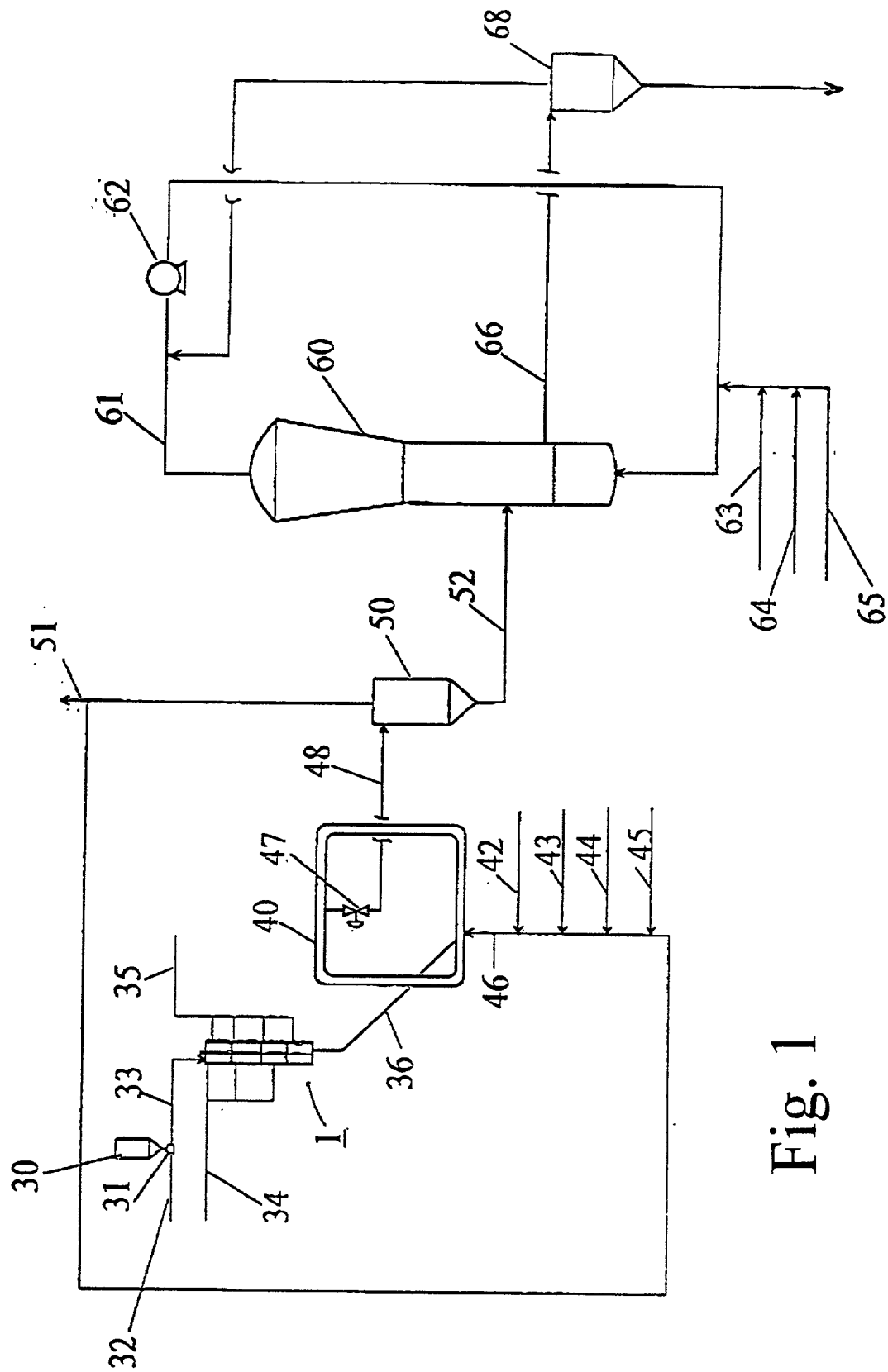
5 40. The process according to claim 39, wherein the catalyst system has been transesterified with a phthalic acid ester - a lower alcohol pair, which transesterifies the catalyst at a temperature of 110 to 150 °C.

10 41. A product prepared by a process according to any of the preceding claims, which is nucleated for higher crystallization temperature, stiffness and optical properties.

42. Use of a homopolymer or copolymer according to any of claims 1 to 4 or a polymer product according to any of claims 5 to 7 for the manufacture of polymer foams, thermoformed and foamed products, films and sheets, and products prepared by blow molding, stretch blow molding, injection moulding or calendering.

(57) Abstract

The present invention concerns a high melt strength propylene polymer or copolymer suitable for manufacturing foams and thermoformed product exhibiting a melt strength of at least 3 g and comprising a high molar mass portion and a low or medium molar mass portion. The polymers are produced by subjecting propylene and optionally other olefins to polymerization in a plurality of polymerization reactors connected in series, employing different amounts of hydrogen as a molar mass modifier in at least two of the reactors, and carrying out the polymerization reaction in the presence of a catalyst system capable of catalyzing the formation of a high molar mass polymerization product having a MFR₂ of less than 0.1 g/10 min and a low or medium molar mass polymerization product having a MFR₂ of more than 0.5 g/10 min.



Fi. 1

3/4

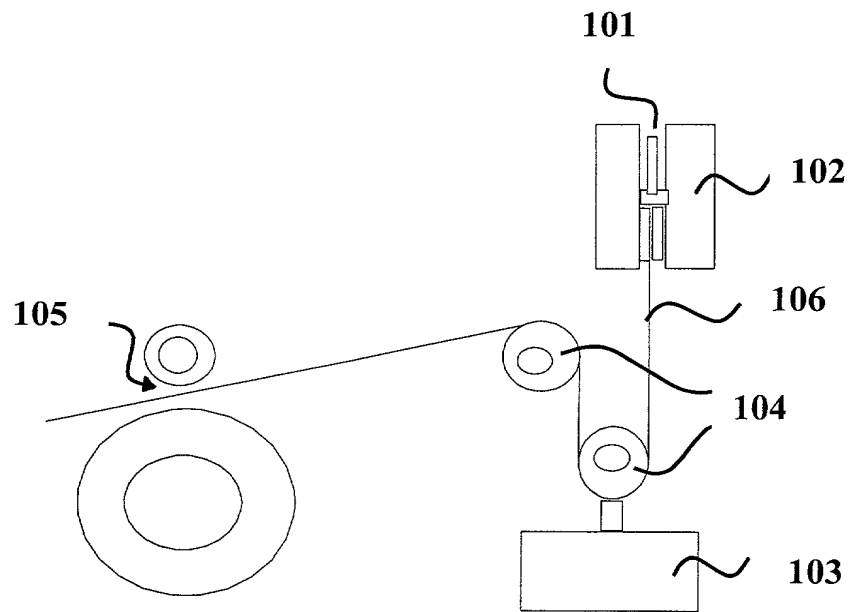
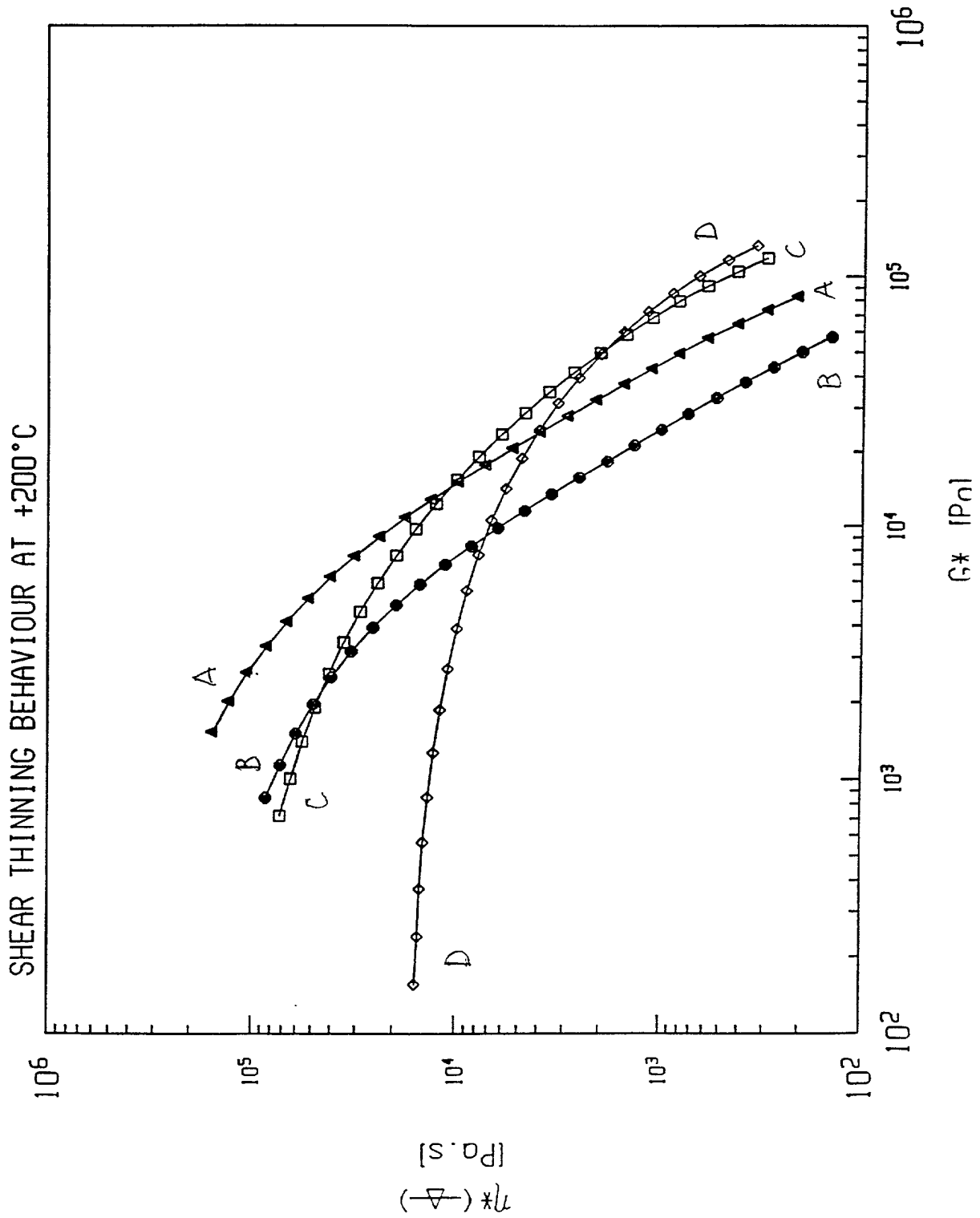


Fig. 3

Fig. 4



BIRCH, STEWART, KOLASCH & BIRCH, LLP

PLEASE NOTE:
YOU MUST
COMPLETE THE
FOLLOWING:

COMBINED DECLARATION AND POWER OF ATTORNEY

ATTORNEY DOCKET NO.

365-442P

FOR PATENT AND DESIGN APPLICATIONS

As a below named inventor, I hereby declare that: my residence, post office address and citizenship are as stated next to my name; that I verily believe that I am the original, first and sole inventor (if only one inventor is named below) or an original, first and joint inventor (if plural inventors are named below) of the subject matter which is claimed and for which a patent is sought on the invention entitled:

Insert Title:

High melt strength polypropylene

Fill in Appropriate
Information -
For Use Without
Specification
Attached:

the specification of which is attached hereto. If not attached hereto,
the specification was filed on March 13, 2000 as
United States Application Number 09/508,405; and /or
the specification was filed on September 28, 1998 as PCT
International Application Number PCT/FI98/00764; and was
amended under PCT Article 19 on _____ (if applicable)

I hereby state that I have reviewed and understand the contents of the above identified specification, including the claims, as amended by any amendment referred to above.

I acknowledge the duty to disclose information which is material to patentability as defined in Title 37, Code of Federal Regulations, §1.56.

I do not know and do not believe the same was ever known or used in the United States of America before my or our invention thereof, or patented or described in any printed publication in any country before my or our invention thereof or more than one year prior to this application, that the same was not in public use or on sale in the United States of America more than one year prior to this application, that the invention has not been patented or made the subject of an inventor's certificate issued before the date of this application in any country foreign to the United States of America on an application filed by me or my legal representatives or assigns more than twelve months (six months for designs) prior to this application, and that no application for patent or inventor's certificate on this invention has been filed in any country foreign to the United States of America prior to this application by me or my legal representatives or assigns, except as follows.

I hereby claim foreign priority benefits under Title 35, United States Code, §119 (a)-(d) of any foreign application(s) for patent or inventor's certificate listed below and have also identified below any foreign application for patent or inventor's certificate having a filing date before that of the application on which priority is claimed:

Insert Priority
Information:
(if appropriate)

Prior Foreign Application(s)

(Number)	(Country)	(Month/Day/Year Filed)	Priority Claimed
973816	Finland	September 26, 1997	<input checked="" type="checkbox"/> Yes <input type="checkbox"/> No
(Number)	(Country)	(Month/Day/Year Filed)	<input type="checkbox"/> Yes <input type="checkbox"/> No
(Number)	(Country)	(Month/Day/Year Filed)	<input type="checkbox"/> Yes <input type="checkbox"/> No
(Number)	(Country)	(Month/Day/Year Filed)	<input type="checkbox"/> Yes <input type="checkbox"/> No
(Number)	(Country)	(Month/Day/Year Filed)	<input type="checkbox"/> Yes <input type="checkbox"/> No

I hereby claim the benefit under Title 35, United States Code, §119(e) of any United States provisional application(s) listed below.

Insert Provisional
Application(s):
(if any)

(Application Number)	(Filing Date)
(Application Number)	(Filing Date)

All Foreign Applications, if any, for any Patent or Inventor's Certificate Filed More Than 12 Months (6 Months for Designs) Prior To The Filing Date of This Application:

Insert Requested
Information:
(if appropriate)

Country	Application No	Date of Filing (Month/Day/Year)

I hereby claim the benefit under Title 35, United States Code, §120 of any United States application(s) listed below and, insofar as the subject matter of each of the claims of this application is not disclosed in the prior United States application in the manner provided by the first paragraph of Title 35, United States Code, §112, I acknowledge the duty to disclose information which is material to patentability as defined in Title 37, Code of Federal Regulations, §1.56 which became available between the filing date of the prior application and the national or PCT international filing date of this application:

Insert Prior U.S.
Application(s):
(if any)

(Application Number)	(Filing Date)	(Status - patented, pending, abandoned)
(Application Number)	(Filing Date)	(Status - patented, pending, abandoned)

I hereby appoint the following attorneys to prosecute this application and/or an international application based on this application and to transact all business in the Patent and Trademark Office connected therewith and in connection with the resulting patent based on instructions received from the entity who first sent the application papers to the attorneys identified below, unless the inventor(s) or assignee provides said attorneys with a written notice to the contrary:

15
Terrell C. Birch (Reg. No. 19,382)
Joseph A. Kolasch (Reg. No. 22,463)
Bernard L. Sweeney (Reg. No. 24,448)
Charles Gorenstein (Reg. No. 29,271)
Leonard R. Svensson (Reg. No. 30,330)
Andrew D. Meikle (Reg. No. 32,868)
Joe McKinney Muncy (Reg. No. 32,334)
C. Joseph Faraci (Reg. No. 32,350)

Raymond C. Stewart (Reg. No. 21,066)
James M. Slattery (Reg. No. 28,380)
Michael K. Mutter (Reg. No. 29,680)
Gerald M. Murphy, Jr. (Reg. No. 28,977)
Terry L. Clark (Reg. No. 32,644)
Marc S. Weiner (Reg. No. 32,181)
Donald J. Daley (Reg. No. 34,313)

Send Correspondence to:

BIRCH, STEWART, KOLASCH & BIRCH, LLP

P.O. Box 747 • Falls Church, Virginia 22040-0747

Telephone: (703) 205-8000 • Facsimile: (703) 205-8050

PLEASE NOTE:
YOU MUST
COMPLETE THE
FOLLOWING:

I hereby declare that all statements made herein of my own knowledge are true and that all statements made on information and belief are believed to be true; and further that these statements were made with the knowledge that willful false statements and the like so made are punishable by fine or imprisonment, or both, under Section 1001 of Title 18 of the United States Code and that such willful false statements may jeopardize the validity of the application or any patent issued thereon.

Full Name of First or Sole
Inventor:
Insert Name of Inventor
Insert Date This
Document is Signed

Insert Residence
Insert Citizenship

Insert Post Office
Address

Full Name of Second
Inventor, if any:

see above

Full Name of Third
Inventor, if any:

see above

Full Name of Fourth
Inventor, if any

see above

Full Name of Fifth
Inventor, if any

see above

GIVEN NAME	FAMILY NAME	INVENTOR'S SIGNATURE	DATE
Päivi	Huovinen	<i>Päivi Huovinen</i>	March 20, 2000 20.3.2000
Residence (City, State & Country)		CITIZENSHIP	
Ferrara, Italy <i>FIX</i>		Finnish	
POST OFFICE ADDRESS (Complete Street Address including City, State & Country)			
Via Ripagrande 63, 44100 Ferrara, Italy			
GIVEN NAME	FAMILY NAME	INVENTOR'S SIGNATURE	DATE
Ali	Harlin	<i>Ali Harlin</i>	Febr. 29, 2000 29.2.2000
Residence (City, State & Country)		CITIZENSHIP	
Vantaa, Finland <i>FIX</i>		Finnish	
POST OFFICE ADDRESS (Complete Street Address including City, State & Country)			
Kärppärinne 1, FIN-01450 Vantaa, Finland			
GIVEN NAME	FAMILY NAME	INVENTOR'S SIGNATURE	DATE
Pirjo	Jääskeläinen	<i>Pirjo Jääskeläinen</i>	Febr. 25, 2000 25.2.2000
Residence (City, State & Country)		CITIZENSHIP	
Porvoo, Finland <i>FIX</i>		Finnish	
POST OFFICE ADDRESS (Complete Street Address including City, State & Country)			
Kaskitie 20, FIN-06150 Porvoo, Finland			
GIVEN NAME	FAMILY NAME	INVENTOR'S SIGNATURE	DATE
Amir	Karbasi	<i>Amir Karbasi</i>	April 8, 2000 8.4.2000
Residence (City, State & Country)		CITIZENSHIP	
Espoo, Finland <i>FIX</i>		Finnish	
POST OFFICE ADDRESS (Complete Street Address including City, State & Country)			
Jousenkaari 11 A 20, FIN-02120 Espoo, Finland			
GIVEN NAME	FAMILY NAME	INVENTOR'S SIGNATURE	DATE
Nina	Manner	<i>Nina Manner</i>	Febr. 28, 2000 28.2.2000
Residence (City, State & Country)		CITIZENSHIP	
Vantaa, Finland <i>FIX</i>		Finnish	
POST OFFICE ADDRESS (Complete Street Address including City, State & Country)			
Keihästie 5 C 53, FIN-01280 Vantaa, Finland			